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Substitute Specification

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TC 2800 MAIL ROOM:

SOLID STATE HETEROJUNCTION AND SOLID STATE SENSITIZED  
PHOTOVOLTAIC CELL.

CROSS-REFERENCE TO RELATED APPLICATIONS.

Not Applicable.

STATEMENT REGARDING FEDERALLY SPONSORED RESEARCH OR  
DEVELOPMENT.

Not Applicable.

INCORPORATION-BY-REFERENCE OF MATERIAL SUBMITTED ON A COMPACT  
DISC.

Not Applicable.

BACKGROUND OF THE INVENTION.

Field of the invention.

The present invention concerns a solid state p-n heterojunction, comprising an electron conductor and a hole conductor. The invention concerns also a solid state sensitized photovoltaic cell, in particular a solar cell.

Description of related art.

The dye sensitized solar cells (DYSC), are more and more maturing into a technically and economically credible alternative to the conventional p-n junction photovoltaics. Photoelectrochemical (PEC) liquid junction cells based on the photosensitization of semiconductor  $\text{TiO}_2$  layers with molecular sensitizers attracted renewed interest after Graetzel et al reported energy conversion efficiencies  $>10\%$ , in Nature, 353 (1991) 737. In 1998, Graetzel et al reported in Nature, 395 (1998) 583, a more innovative all solid state cell based on a heterojunction sensitized by a molecular dye where an amorphous organic hole transport material replaced the liquid electrolyte.

The crucial part in these cells is the dye itself. Only a very limited number of dyes give high photocurrent quantum

yields and are reasonably stable against photo-degradation. Some of the organic dyes exhibiting high light absorption are sensitive to air and water.

It is therefore a target of the present invention to propose photovoltaic cells having improved stability against photo-degradation and environmental influences. It is another target of the invention to achieve higher photocurrent quantum yields, higher photovoltages and hence higher conversion yields in the visible part of the light spectrum.

#### BRIEF SUMMARY OF THE INVENTION.

These aims are achieved by means of a solid state p-n heterojunction comprising a solid electron conductor, - i.e. a n type semiconductor, and a hole conductor, - i.e. a p-type semiconductor, further comprising a sensitising semiconductor, said sensitising semiconductor being located at the interface between the electron conductor and the hole conductor.

For the purpose of sensitization, the sensitizing semiconductor shall be light absorbing in the visible and the near infrared parts of the light spectrum, preferably up to 950 nanometers (eq to 1.4 eV).

The sensitizing semiconductor maybe provided in form of a coating film at the surface of the electron conductor. This coating film is not necessarily continuous. Only a part of the surface of the electron conductor, i.e. a part of the surfaces of the pores of a porous electron conductor may be covered by coating portions.

The sensitizing semiconductor may be provided in form of particles adsorbed at the surface of the electron conductor. The term "particles" herein refers to particles of various sizes, below a nanometer, in the nanometer range, or bigger, but excludes single discrete molecules.

A preferred kind of particles of sensitizing semiconductor for providing a solid state p-n heterojunction according to the present invention are quantum-dots.

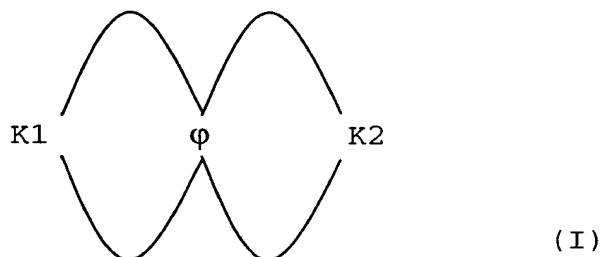
Thereby, a preferred object of the present invention is a new embodiment of a sensitized solar cell where the molecular sensitizer is replaced by a small semiconductor particle, referred to in scientific papers as "nano-dot" or "quantum dot"

or "Q-dot". Quantum dots are semiconductor particles of nanometer size wherein a gradual translation from solid state to molecular structure occurs as the particle size decreases. The quantum dots are adsorbed at the heterojunction constituted by a n-type semiconductor and a p-type semiconductor.

Preferably, the n-type semiconductor is a finely divided and sintered large band gap ceramic oxide. Various metal oxides like  $\text{Ta}_2\text{O}_5$ ,  $\text{Nb}_2\text{O}_5$ ,  $\text{SnO}_2$ ,  $\text{ZnO}$  may be used as solid electronic conductors, whereas nanocrystalline  $\text{TiO}_2$  (anatase) is preferred.

The hole conductor may be selected from hole transporting inorganic solids like copper iodide or copper thiocyanate. Typically, the hole conductor may be an organic charge transport material. This organic charge transport material may be a polymer, like poly-tiophen or poly-arylamin. The hole conductor of the present invention may also be an amorphous, reversibly oxidizable organic non-polymeric compound.

As such a compound, the hole conductor may be selected from the group consisting of spiro- and hetero spiro compounds of general formula (I)



Wherein  $\phi$  is one of C, Si, Ge or Sn, and K1 and K2 are, independently one of the other, conjugated systems. A particularly preferred hole conductor is 2,2',7,7'-tetrakis (N,N-di-p-methoxyphenyl-amine) 9,9'-spirobifluorene, herein further called OMeTAD.

As quantum dots, particles consisting of  $\text{CdS}$ ,  $\text{Bi}_2\text{S}_3$ ,  $\text{Sb}_2\text{S}_3$ , or  $\text{Ag}_2\text{S}$  may be used, whereas  $\text{PbS}$  is preferred. Other compounds suitable for making quantum-dots are In As, **InP**,  $\text{CdTe}$ ,  $\text{CdSe}$ ,  $\text{HgTe}$ . Solid solutions of  $\text{HgTe}$  and  $\text{CdTe}$  or of  $\text{HgSe}$  and  $\text{CdSe}$  are also suitable.

A preferred quantum dot sensitized nanocrystalline heterojunction is constituted of sintered particles of nanocrystalline  $\text{TiO}_2$  onto which PbS particles in the nanometer range are adsorbed as sensitizers, the pores between the particles being filled with amorphous OMeTAD.

The mesoscopic nature of the heterojunction results in a high contact area of the junction, which is prerequisite for efficient solar light harvesting by the quantum dots. Light is absorbed by the Q-dots and produces electron-hole pairs. The electrons are injected from the Q-dots into the electron conducting solid while the holes are injected in the hole conducting side of the junction. In this way electric power is produced from light.

While the sensitization of oxide semiconductors by Q-dots has already been reported for liquid junction cells by Vogel et al, in Chem. Phys. Lett., 174 (1990) 241, these devices show notorious problems with corrosion and photo-corrosion of the Q-dots by the electrolyte, rendering them unsuitable for sensitization. The present invention realises a solid state heterojunction cell that does not present these disadvantages and hence can be used for the conversion of sunlight to electric power.

Quantum dot sensitized heterojunction cells offer several advantages. The band gaps and thereby the absorption ranges are adjustable through the particle size or by adjusting the relative concentrations of components in solid solutions like  $\text{HgTe/CdTe}$  or  $\text{HgSe/CdSe}$ . The band gap of these solutions may be adjusted to approach the optimal value for conversion of sunlight to electric power, which is about 1.4-1.5 eV.

Furthermore, compared to organic dyes, quantum dot sensitization offers improved stability, since the surface of the Q-dots can be modified to improve their photostability.

Another advantage is that the optical cross section of the Q-dots is significantly larger than the one of the molecular dyes. This allows to employ thinner films resulting in higher photovoltages as well as better fill factors of the cell, and hence higher conversion yields. Whereas only a limited number of molecular dyes suit the requirements of DYSC, Q-dot semiconductors have very high extinction coefficients and hence are very effective sensitizers. Since photo generated  $e^-$  and  $h^+$  can be separated effectively as in n-p junctions, quantum dot

sensitization offers a means to combine the mechanical features, efficiency and weather-ruggedness of single crystalline or polycrystalline p-n photovoltaics with an excellent spectral range coverage.

#### BRIEF DESCRIPTION OF THE SEVERAL VIEWS OF THE DRAWINGS.

Other features and advantages of the present invention will appear to those skilled in the art from the following description of an embodiment of a solid state sensitized photovoltaic cell, from the description of a process for making a solid state heterojunction and from measurements made with such a cell, in conjunction with the drawings, among which:

- figure 1 shows a schematic view of a solid state sensitized photovoltaic cell;
- figure 2 is an enlarged view of a heterojunction within the nanocrystalline  $\text{TiO}_2$  layer;
- figure 3 shows absorption spectra of  $\text{TiO}_2$  plates with different deposition treatments; and
- figure 4 shows the photo current generation efficiency as a function of wavelength for Q-dot sensitized solid state heterojunction solar cells.

#### DETAILED DESCRIPTION OF THE INVENTION.

Figure 1 shows the schematic build up of a solid state sensitized photovoltaic cell. From bottom to top there are:

- an ITO transparent glass support 1;
- the glass support 1 is coated by a transparent conducting layer 2 made of F doped  $\text{SnO}_2$ , thereby the coated glass acts as a working electrode, which collects charge and current.
- a dense  $\text{TiO}_2$  layer 3 for avoiding direct contact between the organic hole conductor and the  $\text{SnO}_2$ , which would short circuit the cell;
- a quantum dot sensitized nanocrystalline layer 4, forming a heterojunction, which is build up as will be explained bellow;
- the back contact 5 of the cell is made of a fine gold layer of 10 nanometers covered by a thick nickel layer for a better current collection.

Figure 2 shows a magnified schematic view of the microscopic structure of the layered heterojunction: at the surface of  $\text{TiO}_2$  particles 6, are adsorbed Q-dots 7; the spaces are filled with particles 8 of an organic hole conductor.

Example: process for making a layered heterojunction

The compact  $\text{TiO}_2$  layer is deposited on the  $\text{SnO}_2$ -coated glass by spray pyrolysis. This process is performed following the procedure described in the thesis No. 2187 at EPFL (Ecole Polytechnique Fédérale de Lausanne) of Udo Bach. The nanoporous  $\text{TiO}_2$  layer is deposited by screen printing. The plates are thereafter sintered at a temperature of  $450^\circ\text{C}$  for 15 min. This results in a layer with a thickness of about  $2\text{ }\mu\text{m}$ . The plates are thereafter stored in water free atmosphere.

The technique used to deposit the Q-dots follows the procedure published by Weller et al in J. Phys. Chem. 98, 1994, 3183. Two solutions are needed for this technique, the first one is a saturated  $\text{Pb}(\text{NO}_3)_2$  solution in water, and the second one a  $0.3\text{M}$   $\text{Na}_2\text{S}$  solution in water. The plates are immersed for 1 minute in the first solution. Thereafter, they are rinsed with water and dipped into the second solution for about 10 sec and then rinsed again. This deposition procedure may be repeated several times.

After the deposition of the Q-dots, the hole-conductor is introduced into the pores of the nanocrystalline structure by spin-coating. The spin-coating solution is a 30% by weight solution of OMETAD in chlorobenzene. The OMETAD may be doped with 0.2% of OMETAD<sup>-</sup>. An amount of  $15\mu\text{l}$  of spin-coating solution is used for each plate. Subsequently, the solvent is evaporated.

A semi-transparent gold back contact is evaporated on top of the heterojunction layer under vacuum. A second contact layer made of nickel is sprayed onto the first one.

Absorption spectra are measured with a Hewlett-Packard 8453 spectrometer and are shown in Figure 3. Figure 3 shows that repeating the Q-dot deposition treatment results in an increase of the visible absorption of the electrode. With each deposition, the optical density increases and the electrode becomes strongly coloured. The Q-dots are getting bigger with each treatment.

Figure 4 shows the wavelength response of the photocurrent produced by the device. The incident photon to current conversion efficiency (IPCE) is plotted as a function of the wavelength of the monochromatic light. The conversion yields are high, exceeding 10 percent in the visible. This proves that the sensitization of the heterojunction by the quantum dots functions well.

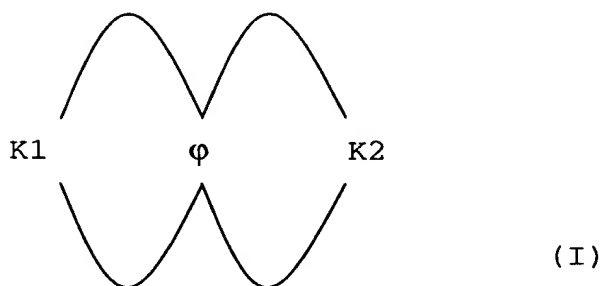
From these measurements, it can be seen that there is an optimum in the number of times the deposition treatment is repeated. For this embodiment, if only one treatment is done, there is not enough Q-dots formed on the  $\text{TiO}_2$  surface. On the other hand, if there are too many deposition treatments, typically 10, as shown in figure 4, the Q-dots get too big and the efficiency drops.

CLAIMS

1. A solid state p-n heterojunction comprising an electron conductor and a hole conductor, characterised in that it further comprises a sensitising semiconductor, said sensitizing semiconductor being located at an interface between said electron conductor and said hole conductor.
2. A heterojunction as claimed in claim 1, wherein said sensitizing semiconductor is in form of a coating film at the surface of said electron conductor.
3. A heterojunction as claimed in claim 1, characterised in that said sensitizing semiconductor is in form of particles adsorbed at the surface of said electron conductor.
4. A heterojunction as claimed in claim 3, characterised in that said sensitizing semiconductor is in form of quantum-dots.
5. A heterojunction as claimed in claim 1, characterised in that said electron conductor is an n-type semiconductor and said hole conductor is a p-type semiconductor.
6. A heterojunction as claimed in claim 5, characterised in that said n-type semiconductor is a ceramic made of finely divided large band gap metal oxide.
7. A heterojunction as claimed in claim 1, characterised in that said electron conductor is nanocrystalline  $\text{TiO}_2$ .
8. A heterojunction as claimed in claim 1, characterised in that said hole conductor is an inorganic hole transporting solid compound.
9. A heterojunction as claimed in claim 1, characterised in that said hole conductor is an amorphous reversibly oxydisable organic or organomethallic compound.
10. A heterojunction as claimed in claim 9, characterised in that said organic compound is a polymer.
11. A heterojunction as claimed in claim 1, characterised in that said hole conductor is selected from the group



consisting of Spiro and Heterospirocompounds of general formula (I),



wherein  $\phi$  is one of C, Si, Ge or Sn, and K1 and K2 are independently one from the other conjugated systems.

12. A heterojunction as claimed in claim 11, wherein said hole conductor is OMeTAD.

13. A heterojunction as claimed in claim 4, wherein said quantum dots are particles consisting of PbS, CdS, Bi<sub>2</sub>S<sub>3</sub>, Sb<sub>2</sub>S<sub>3</sub>, Ag<sub>2</sub>S, InAs, InP, CdTe, CdSe or HgTe or solid solutions of HgTe/CdTe or HgSe/CdSe.

14. A solid state sensitized photovoltaic cell comprising a layered heterojunction as claimed in claim 1.

15. A cell as claimed in claim 14, characterised in that it comprises

- a transparent first electrode,
- a said layered heterojunction and
- a second electrode.

16. A cell as claimed in claim 15, further comprising a dense semiconductive layer between said first electrode and said layered heterojunction.

17. A cell as claimed in claim 14, characterised in that said layered heterojunction is obtained by forming quantum dots on the surface of said electron conductor by at least one deposition treatment, before providing said hole conductor to said layered heterojunction.

18. A cell as claimed in claim 17, characterised in that said deposition treatment is performed 2-10 times before providing said hole conductor.

ABSTRACT OF THE DISCLOSURE

A solid state p-n heterojunction comprising an electron conductor and a hole conductor; it further comprises a sensitizing semiconductor, said sensitizing semiconductor being located at an interface between said electron conductor and said hole conductor. In particular, the sensitizing semiconductor is in form of quantum-dots. A solid state sensitized photovoltaic cell comprises such a layered heterojunction between two electrodes.

SEQUENCE LISTING.

Not Applicable.